DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
Rockville MD 20857

#21

SEP 23 1994

Re: Sporanox®
Docket No. 92E-0472

RECEIVED
SEP 30 PM 4:29
DEPUTY ASSISTANT COMMISSIONER FOR PATENTS

Charles E. Van Horn
Deputy Assistant Commissioner for
Patent Policy and Projects
Office of the Assistant Commissioner for Patents
U.S. Patent and Trademark Office
Crystal Park Building 2, Suite 919
Washington, DC 20231

Dear Mr. Van Horn:

This is in regard to the patent term extension application for U.S. Patent No. 4,267,179 filed by Janssen Pharmaceutica N.V. under 35 U.S.C. § 156. The patent claims the human drug product Sporanox®, NDA 20-083.

In the March 11, 1994 issue of the Federal Register (59 Fed. Reg. 11,612), the Food and Drug Administration published its determination of this product's regulatory review period, as required under 35 U.S.C. § 156(d)(2)(A). The notice provided that on or before September 7, 1994, 180 days after the publication of the determination, any interested person could file a petition with FDA under 35 U.S.C. § 156(d)(2)(B)(i) for a determination of whether the patent term extension applicant acted with due diligence during the regulatory review period.

The 180-day period for filing a due diligence petition pursuant to this notice has expired and FDA has received no such petition. Therefore, FDA considers the regulatory review period determination to be final.

Please let me know if we can provide further assistance.

Sincerely yours,

Ronald L. Wilson
Director
Health Assessment Policy Staff
Office of Health Affairs

cc: Charles J. Metz
Attorney for Applicant
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